

# CR-FREE PAINT COMPOSITIONS AND PAINTED METAL SHEETS

## BACKGROUND OF THE INVENTION

The present invention relates to paint compositions containing a  
5 Cr-free corrosion inhibitor and also a painted metal sheet excellent in corrosion resistance.

A pre-coated steel sheet has been broadly used in various fields such  
as a surfacing member for electric home appliance or an air conditioner and  
10 other goods, due to its good productivity compared with a steel sheet, which is formed to an objective shape in prior to application of paint.

The pre-coated steel sheet is manufactured by applying synthetic resin  
paint to both surfaces or one surface (which will be an external surface of a  
product) of a steel sheet, and baking the applied paint to form a paint layer. A  
15 corrosion inhibitor is commonly added to the resin paint in order to improve corrosion resistance of the paint layer. Adhesiveness of the paint layer is enhanced by chemically converting the surface of a steel sheet to a chromated or phosphated state in prior to paint application. The chemically converted surface layer also effects on improvement of corrosion resistance.

20 In a process of manufacturing a painted steel sheet having a single coat, one kind of paint containing a color pigment and a corrosion inhibitor is applied to the surface and then baked. In a process of manufacturing a painted steel sheet having two or more coats, a primer containing an extender and a corrosion inhibitor is applied to a surface of the steel sheet, and then  
25 intercoat and topcoat paints, which do not contain a corrosion inhibitor, are applied to the undercoat. A back surface of the steel sheet may be also coated with paint containing a corrosion inhibitor.

Chromium compounds or pigments based on chromium compounds,  
e.g. zinc chromate, strontium chromate, red chromate and red silicochromate,  
30 have been used so far as a corrosion inhibitor, due to excellent

corrosion-inhibiting faculty. However, there is a stronger demand in these days for provision of a steel sheet coated with a paint layer free from chromium compounds, accounting harmful influences on the environment. In order to cope with such the demand, a corrosion inhibitor prepared from porous silica particles, to which Ca, Zn, Co, Pb, Sr or Ba ion is bonded by ion-exchange, is proposed instead of chromium compounds.

The proposed corrosion inhibitor captures corrosive ions such as hydrogen ion and discharges the bonded ion in return. Especially, Ca ion-bonded corrosion inhibitor performs good corrosion-inhibiting faculty. Such the corrosion inhibitor is ordinarily added to paint at a ratio of 2-50 parts by weight based on 100 parts by weight of resinous components in a paint layer. However, the Ca ion-bonded corrosion inhibitor is somewhat insufficient of corrosion and moisture resistance in comparison with chromium compounds, so that blisters often occur in a paint layer in a wet atmosphere.

#### SUMMARY OF THE INVENTION

The present invention aims at provision of a metal sheet coated with a resin paint containing a corrosion inhibitor prepared from Ca ion-bonded silica particles whose corrosion inhibiting faculty is enhanced by coexistence of a polyphosphate.

The present invention proposes new paint composition containing a corrosion inhibitor prepared from porous silica particles, to which Ca ion is bonded by ion-exchange, together with a polyphosphate at a ratio predetermined in relation with a ratio of the corrosion inhibitor.

The polyphosphate is preferably aluminum pyrophosphate, aluminum metaphosphate or aluminum dihydrogentriphosphate. Especially, aluminum dihydrogentriphosphate performs best results. The polyphosphate may be used in a state mixed or coated with zinc oxide, titanium oxide, magnesium compound, a silane coupling agent or silicone oil,

to improve corrosion resistance and storage stability.

Ca-bonded silica particles (hereinafter referred to as "the corrosion inhibitor A") are generally added to resin paint at a ratio of 2-50 parts by weight based on 100 parts by weight of resinous components of the paint. The polyphosphate (hereinafter referred to as "the polyphosphate B") is added to the resin paint at an A/B weight ratio of 60/40 to 5/95 and at an A+B ratio of 5-150 parts by weight based on 100 parts by weight of resinous components of the paint.

The proposed paint is used for formation of an undercoat or a single coat on a base metal sheet such as a steel sheet plated with a Zn, alloyed Zn, Zn-5%Al or Zn-55%Al layer. A surface of the base metal sheet is preferably chemically processed to a state affinitive to the paint.

#### DETAILED DESCRIPTION OF THE PREFERRED EMBODIMENTS

The inventors have researched the reason why the corrosion inhibitor A does not exhibit well corrosion inhibiting faculty as compared with a conventional corrosion inhibitor based on chromium compounds especially in a wet atmosphere, and discovered that an effect of Ca ion bonded to silica particles is not maintained over a long term since Ca ion is easy to dissolve in water permeated into a paint layer.

The corrosion inhibitor A could be improved in moisture resistance by coating silica particles with hydrophobic films such as a silane coupling agent or silicone oil in order to suppress dissolution of Ca ion. However, such the hydrophobic coating closes pores of the silica particles and scarcely permits dissolution of Ca ion, resulting in degradation of corrosion resistance. The inventors have researched and examined various means to suppress dissolution of Ca ion, and hit upon addition of a polyphosphate to paint.

The polyphosphate B makes ionic bond similar to chelate bond on surfaces of silica particles. The ionic bond controls dissolution of Ca ion, but does not completely inhibit dissolution of Ca ion. Dissolution of Ca ion may be

also suppressed by a pH-buffering effect of the polyphosphate B to moderate acidification caused by corrosive ions such as hydrogen ion.

The polyphosphate B is preferably added to resin paint at an A/B weight ratio of 60/40 to 5/95 and also at an A+B ratio of 5-150 parts by weight based on 100 parts by weight of resinous components of the paint, in order to most efficiently realize the effect of the polyphosphate B.

Addition of the polyphosphate B at the controlled ratio ensures continuous dissolution of Ca ion over a long term without degradation of the corrosion inhibitor A. Excessive addition of the polyphosphate B unfavorably suppresses dissolution of Ca ion, while shortage of the polyphosphate B causes occurrence of blisters in a paint layer due to poor moisture resistance. The A+B ratio in the range of 5-150 parts by weight is also important to form a paint layer good of corrosion resistance, adhesiveness and workability.

Ca ion is bonded to silica particles as the corrosion inhibitor A at a ratio of 3-40% in general. A ratio of the polyphosphate B is adjusted in response to amounts of the corrosion inhibitor A. For instance, a ratio of the polyphosphate is kept higher as increase of Ca ion bonded to the corrosion inhibitor A, but kept lower as decrease of Ca ions bonded to the corrosion inhibitor A.

There are no restrictions on a kind of resin paint, to which the corrosion inhibitor A and the polyphosphate B are added, but ordinarily based on polyester, macromolecular polyester, epoxy, epoxy-denatured polyester, epoxy-denatured macromolecular polyester or polyether sulfonate. Molecular weight, glass transition temperature and cross-link density of the resin paint are properly adjusted together with ratios of a curing agent and other pigments, accounting use of a painted steel sheet. For instance, a paint layer good of workability is obtained from resin paint, whose glass transition temperature is adjusted at a level of 40°C or lower to increase elongation at a higher ratio above 50%.

The resin paint may be applied as a single coat on a metal sheet or as



water. Metal components originated in the fluoroacid are reacted with hydroxyl groups of resin primer and also reacted with phosphoric acid discharged from the polyphosphate B in the undercoat. The surface layer of the metal sheet, which has been activated by etching, is reacted with Ca ion  
5 supplied from the corrosion inhibitor A in the undercoat, so bonding reaction of resin paint to the surface is accelerated. Consequently, the undercoat is firmly fixed to the surface of the base metal sheet, and a corrosion-resistant layer is generated at the boundary between the base metal sheet and the undercoat. Fluoric ion is also reacted with Ca ion supplied from the corrosion  
10 inhibitor A in the undercoat and converted to a stable compound such as calcium fluoride. The stable compound serves as a barrier and further strengthens the fluorided surface layer.

The fluorided coat is preferably formed at a ratio of 0.5-500mg/m<sup>2</sup> calculated as deposited fluorine or at a ratio of 0.1-500mg/m<sup>2</sup> calculated as  
15 deposited metals such as Ti, Zr, Hf, Si, Ge, Sn and B in total, in order to gain most advantageous effects on corrosion resistance.

The fluorided coat may further contain one or more of oxides, hydroxides, phosphates, carbonates or other salts of Ti, Zr, Mo, Si, Al, Hf, Ge and Sn. These compounds are dissolved or dispersed in the chemical agent.  
20 Other additives, e.g. polyvinyl alcohol, tannic acid, starch, corn starch, water-soluble or dispersible polymer such as acrylic resin, a silane coupling agent, a titanium coupling agent, a surfactant, etc., may be optionally added to the chemical agent.

The chemical agent is applied to a surface of a base metal sheet by a  
25 roll-type applicator, spraying or dipping. Thereafter, the surface of the base metal sheet is dried without washing, to form the fluorided coat thereon.

After the fluorided coat is formed on a surface of a base metal sheet, resin paint is applied to the surface by a conventional paint-applying process such as roll-type applying, curtain-flow applying, spraying or dipping. The  
30 applied paint is then baked to form an undercoat or a single coat on the

surface of the base metal sheet. Primer paint may contain various additives, e.g. inorganic pigments such as titanium oxide, calcium carbonate and silica other than the corrosion inhibitor A and the polyphosphate B.

Various resin paint such as polyester, urethane, vinyl chloride, acrylic, polyether sulfonate, silicone or fluoric resin may be used for formation of a topcoat without any special restrictions. A kind, molecular weight, glass-transition temperature and pigmentary dosage of topcoat resin paint are properly adjusted in response to use of a painted metal sheet as an interior member, cladding, etc.. PTFE (polytetrafluoroethylene) may be added to the topcoat paint. An intercoat may be optionally formed on an undercoat in prior to application of topcoat paint, by 3-coat process. Of course, the painted metal having a single coat can be manufactured by 1-coat process. Paint applied to a base metal sheet is ordinarily baked by hot air in a continuous coating line.

#### EXAMPLE

##### Example 1:

Galvanized steel sheets (of 0.5mm in thickness, Zn deposition ratio of 45g/m<sup>2</sup> per single surface) were chemically pretreated with a Cr-free phosphating agent (I), II a Cr-free silicating agent (II), a Cr-free zirconating agent (III), a Cr-free manganating agent (IV), and a Cr-free titanating agent (V), as shown in Table 1.

Thereafter, epoxy-denatured macromolecular polyester resin paint was applied to each steel sheet and baked 40 seconds at a maximum temperature of 215°C to form a paint layer of 10μm in dry thickness.

Paint compositions used in Example 1 contained the corrosion inhibitor A (i.e., porous silica particles to which Ca ion was bonded by ion-exchange) and the polyphosphate B. Ratios of the polyphosphate in relation with the corrosion inhibitor A are shown in Table 1.

Table 1: PAINT COMPOSITIONS USED IN EXAMPLE 1

NOTE	No	pretreatment	a polyphosphate B			a total ratio (parts by weight) of A+B	color tone
			kind	a ratio of B (parts by weight)	a weight ratio of A/B		
INVENTIVE EXAMPLES	1	I	a	5	10/90	20	white
	2	I	a	10	10/90	40	white
	3	I	a	15	10/90	60	white
	4	I	a	5	10/90	50	white
	5	I	a	13	25/75	50	white
	6	I	a	25	50/50	50	white
	7	II	a	13	25/75	50	white
	8	III	a	13	25/75	50	white
	9	IV	b	13	25/75	50	white
	10	V	a	13	25/75	50	white
	11	I	b	13	25/75	50	white
	12	I	c	13	25/75	50	white
COMPARATIVE EXAMPLES	1	I	a	0.5	25/75	2	white
	2	I	a	0.5	5/95	10	white
	3	I	a	40	25/75	160	white
	4	I	a	0	0/100	50	white
	5	I	a	1.5	3/97	50	white
	6	I	a	40	80/20	50	white
	7	I	a	50	100/0	50	white
	8	I	a	10	5/95	200	white
	9	I	zinc phosphate			50	white
	10	I	wet silica			50	white
	11	I	wet silica/a=25/75			50	white
	12	I	silica particles/zinc phosphate=25/75			50	white
	13	I	strontium chromate			10	yellow



Notes for Table 1

- 1: A polyphosphate a is aluminum dihydrogentripolyphosphate, b is aluminum metapolyphosphate, and c is aluminum pyrophosphate.
- 2: A total ratio of the corrosion inhibitor A and the polyphosphate B is based on 100 parts by weight of resinous components in a paint layer.
- 3: Color tone white is derived from titanium oxide added to paint, and yellow is derived from strontium chromate added to paint.

After the applied paint was dried, each painted steel sheet was examined by the following coating performance test. Test results are shown in Table 2 (for inventive examples) and Table 3 (for comparative examples).

Test dipping in boiling water

Each test piece was immersed 2 hours in boiling water and raised therefrom. The test piece was observed at its external appearance and evaluated as follows:

- : no faults
- △ : occurrence of blisters or dull appearance
- × : occurrence of great blisters or dull appearance

The test piece was then examined by a taping test, whereby the test piece was bent with 0t, an adhesive tape was stuck onto a bent part, and then the adhesive tape was peeled off the bent part. Thereafter, the paint layer remaining on the surface of the steel sheet was checked and evaluated as follows:

- : no faults
- △ : partial peeling-off of a paint layer
- × : significant peeling-off of a paint layer

Tests for corrosion and moisture resistance

Each test piece was cross-cut with injuries extending to a steel base, and then subjected to a 240 hours-salt water spray test regulated by JIS Z2371 as a corrosion resistance test. The same cross-cut test piece was held

240 hours in an atmosphere at 50°C with 98%RH for a moisture resistance test. Thereafter, the test piece was observed at its flat part, and maximum widths of blisters in a paint layer were measured at a lower burred part of a side face and at one side of a cross-cut part. Results were evaluated as

5 follows:

External appearance at a flat part

○ : no faults

△ : occurrence of blisters or dull surface

× : significant blisters or dull surface

10 Maximum width of blisters at a lower burred part of a side face

◎ : not more than 1mm

○ : 1-3mm

△ : 3-6mm

× : more than 6mm

15 Maximum width of blisters at one side of a cross-cut part

◎ : no blisters

○ : not more than 1mm

△ : 1-2mm

× : more than 2mm

Table 2: PROPERTIES OF PAINTED STEEL SHEETS (INVENTIVE EXAMPLES)

No	resistance to boiling water		corrosion resistance			moisture resistance		
	appearance	taping test	appearance at a flat part	width of blisters at a lower burred part of a side face	width of blisters at cross-cut part	appearance at a flat part	width of blisters at a lower burred part of a side face	width of blisters at cross-cut part
1	○	○	○	○	◎	○	◎	◎
2	○	○	○	◎	◎	○	◎	◎
3	○	○	○	◎	◎	○	◎	◎
4	○	○	○	○	◎	○	◎	◎
5	○	○	○	◎	◎	○	◎	◎
6	○	○	○	◎	◎	○	◎	◎
7	○	○	○	◎	◎	○	◎	◎
8	○	○	○	◎	◎	○	◎	◎
9	○	○	○	◎	◎	○	◎	◎
10	○	○	○	◎	◎	○	◎	◎
11	○	○	○	◎	◎	○	◎	◎
12	○	○	○	◎	◎	○	◎	◎

Table 3: PROPERTIES OF PAINTED STEEL SHEETS (COMPARATIVE EXAMPLES)

No	resistance to boiling water		corrosion resistance			moisture resistance		
	appearance	taping test	appearance at a flat part	width of blisters at a lower burred part of a side face	width of blisters at cross-cut part	appearance at a flat part	width of blisters at a lower burred part of a side face	width of blisters at cross-cut part
1	○	○	○	×	×	○	○	○
2	○	○	○	×	×	○	○	○
3	△	△	○	⊙	⊙	△	⊙	⊙
4	○	○	○	×	×	○	⊙	⊙
5	○	○	○	△	△	○	⊙	⊙
6	△	○	○	⊙	⊙	△	⊙	⊙
7	×	△	○	⊙	⊙	△	⊙	⊙
8	○	△	○	○	○	○	⊙	⊙
9	○	○	○	×	×	○	⊙	⊙
10	○	○	○	×	×	○	⊙	⊙
11	○	○	○	△	△	○	⊙	⊙
12	△	○	○	○	○	○	⊙	⊙
13	○	○	○	○	○	○	⊙	⊙

Example 2:

After the same steel sheets as in Example 1 were chemically pretreated in the same way, epoxy-denatured macromolecular polyester primer paint was applied to each steel sheet and then baked 30 seconds at a maximum temperature of 215°C to form an undercoat of 5μm in dry thickness. Thereafter, macromolecular polyester topcoat paint was applied to the steel sheet and baked 40seconds at a maximum temperature of 230°C to form a topcoat film of 15μm in dry thickness.

Primer paints used in Example 2 had compositions containing the corrosion inhibitor A and the polyphosphate B, as shown in Table 4, while other primer paints for comparison had compositions shown in Table 5.

Properties of paint layers examined by the same tests as in Example 1 are shown in Table 6 (for inventive examples) and Fig. 7 (for comparative examples).

Table 4: PRIMER PAINT COMPOSITIONS (INVENTIVE EXAMPLE)

No	pretreatment	polyphosphate B			a total ratio (parts by weight) of A+B
		kind	a ratio of B (parts by wt.)	a weight ratio of A/B	
1	I	a	2	25/75	8
2	I	a	5	25/75	20
3	I	a	10	25/75	40
4	I	a	15	25/75	60
5	I	a	20	25/75	80
6	I	a	30	25/75	120
7	I	a	5	10/90	50
8	I	a	13	25/75	50
9	I	a	25	50/50	50
10	II	a	13	25/75	50
11	III	a	13	25/75	50
12	IV	a	13	25/75	50
13	V	a	13	25/75	50
14	I	a	13	25/75	50
15	I	c	13	25/75	50

Table 5: PRIMER PAINTS COMPOSITIONS (COMPARATIVE EXAMPLES)

No	pretreatment	polyphosphate B			a total ratio (parts by weight) of A+B
		kind	a ratio of B (parts by wt.)	a weight ratio of A/B	
16	I	a	0.5	25/75	2
17	I	a	0.5	5/95	10
18	I	a	60	25/75	240
19	I	a	0	0/100	50
20	I	a	1.5	3/97	50
21	I	a	40	80/20	50
22	I	a	50	100/0	50
23	I	a	10	5/95	200
24	I	zinc phosphate			50
25	I	wet silica			50
26	I	wet silica/ a =25/75			50
27	I	silica particles/zinc phosphate=25/75			50

Table 6: PROPERTIES OF PAINT LAYERS (INVENTIVE EXAMPLES)

TABLE OF PROPERTIES OF PAINTS (CONVENTIONAL EXAMPLES)									
No	resistance to boiling water		corrosion resistance			moisture resistance			width of blisters at a cross-cut part
	appearance	taping test	appearance at a flat part	width of blisters at a lower burred part of a side face	width of blisters at a cross-cut part	appearance at a flat part	width of blisters at a lower burred part of a side face		
1	○	○	○	○	○	○	○	○	○
2	○	○	○	◎	◎	○	◎	◎	◎
3	○	○	○	◎	◎	○	◎	◎	◎
4	○	○	○	◎	◎	○	◎	◎	◎
5	○	○	○	◎	◎	○	◎	◎	◎
6	○	○	○	◎	◎	○	◎	◎	◎
7	○	○	○	◎	◎	○	◎	◎	◎
8	○	○	○	◎	◎	○	◎	◎	◎
9	○	○	○	◎	◎	○	◎	◎	◎
10	○	○	○	◎	◎	○	◎	◎	◎
11	○	○	○	◎	◎	○	◎	◎	◎
12	○	○	○	◎	◎	○	◎	◎	◎
13	○	○	○	◎	◎	○	◎	◎	◎
14	○	○	○	◎	◎	○	◎	◎	◎
15	○	○	○	◎	◎	○	◎	◎	◎

Table 7: PROPERTIES OF PAINT LAYERS (COMPARATIVE EXAMPLES)

No	resistance to boiling water		corrosion resistance			moisture resistance		
	appearance	taping test	appearance at a flat part	width of blisters at a lower burred part of a side face	width of blisters at a cross-cut part	appearance at a flat part	width of blisters at a lower burred part of a side face	width of blisters at a cross-cut part
16	○	○	○	△	△	○	○	○
17	○	○	○	△	△	○	○	○
18	○	△	○	◎	◎	○	◎	◎
19	○	○	○	△	△	○	◎	◎
20	○	○	○	△	△	○	◎	◎
21	△	○	○	◎	◎	△	◎	◎
22	×	△	○	◎	◎	△	◎	◎
23	○	△	○	○	○	○	◎	◎
24	○	○	○	×	×	○	◎	◎
25	○	○	○	×	×	○	◎	◎
26	○	○	○	△	△	○	◎	◎
27	△	○	○	○	○	○	◎	◎



Example 3:

[Example]

Various plated steel sheets of 0.5mm in thickness were surface-conditioned with a ratio of  $10\text{mg/m}^2$  calculated as deposited Ni or  
5 alkali-degreased. A chemical agent having composition shown in Table 8 was applied to each steel sheet by a bar-type applicator at a ratio of  $1\text{ ml/m}^2$  and dried at  $100^\circ\text{C}$  as pre-treatment for application of paint.

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Table 8: COMPOSITIONS OF CHEMICAL AGENTS FOR PRE-TREATMENT OF BASE STEEL SHEETS

	inventive examples								comparative examples				
Agent No.	1	2	3	4	5	6	7	8	9	10	11	12	13
fluorotitanate	65.20				48.60	50.40	0.78						
Fluorozirconate		35.20						5.59					
Fulurosilicate			35.20										
Fluoroborate				35.20									
deionized water	920.50	920.50	920.50	920.50	826.70	869.40	998.00	760.74	920.5	826.70	869.40	998.00	760.74
colloidal silica	5.90	5.90	5.90	5.90		5.00			5.90		5.00		
vaporized silica	3.90												
zirconium hydroxide	4.50	10.50	10.50	10.50					10.50				
zirconium carbonate					10.00	15.00				10.00	15.00		
molybdenum hydroxide								113.20					113.20
manganese oxide					20.00					20.00			
hydrogen fluoride							0.10						
phosphoric acid					34.50		0.67	106.30		34.50		0.67	106.30
tannic acid	5.00												
Starch								14.17					14.17
polyvinyl alcohol	5.00												
aminomethylated polyvinyl phenol		27.90	27.90	27.90	61.20	61.20	0.45		27.90	61.20	61.20	0.45	

A processed surface layer was examined by X-ray fluorescence analysis to measure a total amount of F, Ti, Zr, Hf, Si, Ge, Sn and B deposited thereon. Results are shown in Table 9.

5 Table 9: DEPOSITION RATIOS (by mg/m<sup>2</sup>) IN RESPONSE TO  
KINDS OF OF CHEMICAL AGENTS

Inventive Examples	Agent No.	1	2	3	4	5	6	7	8
	fluorides	40.9	17.0	31.1	32.6	39.1	34.7	0.6	2.9
	metals	35.8	28.5	16.2	13.9	19.5	23.2	0.3	2.6
Comparative Examples	Agent No.	9	10	11	12	13			
	fluorides	0	0	0	5.8	0			
	metals	10.5	5.3	12.6	0	0			

A ratio of metals means a total ratio of Ti, Zr, Hf, Si, Ge Sn and B deposited on a surface of a base steel sheet

Each base steel sheet was processed with a chemical agent to form a fluorided layer at a ratio calculated as deposited fluorine, as shown in Table  
10 10 (for inventive examples) and Table 11 (for comparative examples).

Table 10: PREPARATION OF STEEL SHEETS FOR PAINT APPLICATION  
(INVENTIVE EXAMPLES)

No.	a coated-base steel sheet		a fluoride coat	
	a plating layer	an adhesion ratio (g/m <sup>2</sup> )	a chemical agent No.	an adhesion ratio(mg/ m <sup>2</sup> )
1	Zn	45	1	50
2	Zn	45	1	50
3	Zn	45	1	50
4	Zn	45	1	50
5	Zn	45	1	50
6	Zn	45	1	10
7	Zn	45	1	100
8	Zn	45	2	50
9	Zn	45	3	50
10	Zn	45	4	50
11	Zn	45	5	50
12	Zn	45	6	50
13	Zn	45	7	50
14	Zn	45	8	50
15	Zn-5%Al	45	1	50
16	Zn-55%Al	70	1	50
17	Al	70	1	50
18	Zn-6Al-3%Mg	70	1	50
19	Zn-55%Al	70	1	50
20	Zn-55%Al	70	7	50

Table 11: PREPARATION OF STEEL SHEETS FOR PAINT APPLICATION  
(COMPARATIVE EXAMPLES)

No.	a coated-base steel sheet		a fluoride ciat	
	a plating layer	an adhesion ratio (g/m <sup>2</sup> )	a chemical agent No.	an adhesion ratio (mg/m <sup>2</sup> )
1	Zn	45	1	50
2	Zn	45	1	50
3	Zn	45	1	1
4	Zn	45	9	50
5	Zn	45	10	50
6	Zn	45	11	50
7	Zn	45	12	50
8	Zn	45	13	50
9	Zn	45	phosphating	
10	Zn	45	tannic acid	

After the chemical processing, two types of painted steel sheets were  
5 manufactured as follows:

The first group of painted steel sheets are coated with undercoats of 5µm in dry thickness and topcoats of 15µm in dry thickness. Primer paints based on a thermosetting epoxy-denatured macromolecular polyester were applied to the surface of the steel sheet and baked at a maximum temperature of 215°C, while macromolecular polyester topcoat paints were applied to the undercoats and baked 40seconds at a maximum temperature of 230°C.

The second group of painted steel sheets were coated with undercoats of 5µm in dry thickness and topcoats of 15µm in dry thickness. Primer paints based on a polyether sulfonate were applied to the steel sheets and baked 120 seconds at a maximum temperature of 340°C, while topcoat

paints based on a polyether sulfonate containing PTFE (polytetrafluoroethylene) were applied to the undercoats and baked 180 seconds at a maximum temperature of 410°C.

In each case, paint compositions (shown in Tables 12 and 13) containing the corrosion inhibitor A and the polyphosphate B were used as primer paints.

In some of comparative examples, painted steel sheets were prepared from base steel sheet which were subjected to conventional chemical converting treatment using a phosphate or tannic acid, or to chemical converting treatment using the same chemical agent as in the inventive examples except for omission of fluoroacids. In others of comparative examples, primer paints containing either one of the corrosion inhibitor A and the polyphosphate B were also applied to base steel sheets.

Table 12: PRIMER PAINT COMPOSITIONS (INVENTIVE EXAMPLES)

No.	base resin	a corrosion inhibitor A (parts by wt.)	an A/B weight ratio	an A+B ratio (parts by wt.)
1	I	2	25/75	8
2	I	13	25/75	50
3	I	30	25/75	150
4	I	5	10/90	50
5	I	25	50/50	50
6	I	13	25/75	50
7	I	13	25/75	50
8	I	13	25/75	50
9	I	13	25/75	50
10	I	13	25/75	50
11	I	13	25/75	50
12	I	13	25/75	50
13	I	13	25/75	50
14	I	13	25/75	50
15	I	13	25/75	50
16	I	13	25/75	50
17	I	13	25/75	50
18	I	13	25/75	50
19	II	13	25/75	50
20	II	13	25/75	50

Base resin I is epoxy-denatured macromolecular polyester primer with macromolecular polyester topcoat, and II is polyether sulfonate (PES) primer with PES/PTFE topcoat.

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Table 13: PRIMER PAINT COMPOSITIONS (COMPARATIVE EXAMPLES)

No.	base resin	a corrosion inhibitor A (parts by wt.)	an A/B weight ratio	an A+B ratio (parts by wt.)
1	I	ion-exchanged silica solely		50
2	I	aluminum polyphosphate solely		50
3	I	13	25/75	50
4	I	13	25/75	50
5	I	13	25/75	50
6	I	13	25/75	50
7	I	13	25/75	50
8	I	13	25/75	50
9	I	13	25/75	50
10	I	13	25/75	50

Each painted steel was examined by the same tests as in Example 1 to research properties of a paint layer such as boiling water resistance, corrosion resistance and moisture resistance. Results are shown in Tables 14 (for inventive examples) and Table 15 (for comparative examples).



Table 14: PROPERTIES OF PAINT LAYERS (INVENTIVE EXAMPLES)

No.	Resistance to boiling water		corrosion resistance			moisture resistance		
	appearance	Taping test	at a flat part	at a lower burred part	at a cross -cut part	at a flat part	at a lower burred parts	at a cross -cut part
1	○	○	○	○	○	○	○	○
2	○	○	○	○	○	○	○	○
3	○	○	○	○	○	○	○	○
4	○	○	○	○	○	○	○	○
5	○	○	○	○	○	○	○	○
6	○	○	○	○	○	○	○	○
7	○	○	○	○	○	○	○	○
8	○	○	○	○	○	○	○	○
9	○	○	○	○	○	○	○	○
10	○	○	○	○	○	○	○	○
11	○	○	○	○	○	○	○	○
12	○	○	○	○	○	○	○	○
13	○	○	○	○	○	○	○	○
14	○	○	○	○	○	○	○	○
15	○	○	○	○	○	○	○	○
16	○	○	○	○	○	○	○	○
17	○	○	○	○	○	○	○	○
18	○	○	○	○	○	○	○	○
19	○	○	○	○	○	○	○	○
20	○	○	○	○	○	○	○	○

Table 15: PROPERTIES OF PAINT LAYERS (COMPARATIVE EXAMPLES)

No.	Resistance to boiling water		corrosion resistance			moisture resistance		
	appearance	taping test	at a flat part	at a lower burred part	at a cross cut part	at a flat part	at a lower burred part	at a cross cut part
1	×	△	○	○	○	△	○	○
2	○	○	○	△	△	○	○	○
3	○	△	○	△	△	○	⊙	⊙
4	○	△	○	△	△	○	⊙	⊙
5	○	△	○	△	△	○	⊙	⊙
6	○	△	○	△	△	○	⊙	⊙
7	○	△	○	△	△	○	⊙	⊙
8	○	△	○	△	△	○	⊙	⊙
9	○	○	○	△	△	○	⊙	⊙
10	○	○	○	×	×	○	⊙	⊙

It is noted from Table 15 that paint layers formed on steel sheets as comparative examples had disadvantages on at least one or more of boiling water resistance, corrosion resistance and moisture resistance. On the other hand, paint layers (shown in Table 14) formed on steel sheets as inventive  
5 examples were superior any of boiling water resistance, corrosion resistance and moisture resistance. The comparison proves that painted steel sheets according to the present invention are excellent in all of boiling water resistance, corrosion resistance and moisture resistance without inclusion of chromium compounds.

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According to the present invention as above-mentioned, there is provided a painted steel sheet excellent in corrosion and moisture resistance due to co-existence of the corrosion inhibitor A and the polyphosphate B in a paint layer without use of chromium compounds, which would put harmful  
15 influences on the environment. Since the polyphosphate B controls dissolution of Ca ion from the corrosion inhibitor A, the corrosion inhibitor A maintains its corrosion-inhibiting faculty over a long term.